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(11) **EP 0 528 988 B1**

(12) **EUROPEAN PATENT SPECIFICATION**

(45) Date of publication and mention
of the grant of the patent:
16.04.1997 Bulletin 1997/16

(51) Int Cl.⁶: **G01N 33/552**

(86) International application number:
PCT/US91/03311

(21) Application number: **91911210.2**

(87) International publication number:
WO 91/18292 (28.11.1991 Gazette 1991/27)

(22) Date of filing: **16.05.1991**

(54) **HIGHLY REFLECTIVE BIOGRATINGS AND METHOD**

HOCHREFLEKTIVE BIOGITTER UND VERFAHREN

RESEAUX DE DIFFRACTION BIOLOGIQUES HAUTEMENT REFLECHISSANTS ET PROCEDE ASSOCIE

(84) Designated Contracting States:
DE ES FR GB IT

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(30) Priority: **17.05.1990 US 525828**

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(43) Date of publication of application:
03.03.1993 Bulletin 1993/09

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EP-A- 0 178 083 **US-A- 4 647 544**
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• **PATENT ABSTRACTS OF JAPAN vol. 012, no.**
307 (P-747)22 August 1988 & JP-A-63 078 051
(TEIJIN LTD) 8 April 1988

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Description

This invention relates to improved devices for use in a reflective diffraction immunoassay and their method of manufacture. In particular, this invention relates to multilayer biogratings having high reflectivity and high protein binding capacity.

Many solid-phase binding assays involve surface illumination and consequent light emissions from molecules attached to the surface or are masked by forward scattering. Generally, these emissions travel in all directions. Either these divergent emissions must be collected with expensive and awkward light collection optics to achieve sensitivity, the inherent inefficiencies and consequent low signal to light level ratios must be accepted, or the signal must be measured against a strong background.

Diffraction gratings cause light to be diffracted into specific angles as contrasted to being scattered in all directions. The original diffraction gratings were prepared by ruling a number of straight, parallel grooves in a surface. Incident light was diffracted by each of the surfaces and was principally directed in directions in which light from each groove interferes constructively with light scattered by the other grooves. This constructive light interference property of a grating allows efficient collection of light. Preformed diffraction gratings of this type have been used in binding assay systems.

Many assay systems have been developed using different physically measurable properties of reagents to provide a measurement of an analyte concentration in a sample. Radioimmunoassay (RIA), immunofluorescence, chemiluminescence, enzyme immunoassays (EIA), free radical immunoassays (FRAT), light scattering nephelometry, transistor bridge probes, indium reflective surfaces, and ultrasonic probes have been applied. These systems use the highly selective reaction between a primary member of a binding pair such as an antibody or antigen and an analyte selectively binding therewith. These techniques require expensive measurement equipment and often involve very complicated test procedures.

Reflective and transmissive biograting immunoassay systems and methods were disclosed in U.S. Patent 4,647,544. One embodiment described in the patent uses a biograting, a substantially flat surface having a coating thereon and having substantially uniform light scattering properties. The coating comprises a diffraction grating pattern of alternating parallel linear zones of an active and deactivated binding reagent. The zones form a diffraction grating when the active binding reagent binds with its opposite member of the binding pair. In the absence of such binding, no significant light diffraction occurs, that is, light energy detected at the diffraction angles is at a minimum value, approaching zero. When the binding occurs, the accumulation of bound material in the patterns of a diffraction grating creates a light disturbing grating, and light detected at the

light diffraction angles increases to a larger value which correlates to the presence and quantity of the binding partner (analyte) in the sample. The flat surfaces upon which the biograting is formed in the patent include glass, plastic, plastic coating on a solid surface, gel or other suitable inert material onto which specific antibody molecules can be attached.

U.S. Patent 4,876,208 describes transmissive and reflective diffraction binding assays and biograting systems of the type described in U.S. Patent 4,647,544. The biograting supports disclosed in this patent include a smooth upper surface of any material to which a primary hybridizing reagent can be adhered by physical or chemical bonding and which will not interfere with the reactions which are used to determine the presence and extent of the hybridizing reaction. Organic and inorganic polymers, both natural and synthetic, are described. Examples of polymers listed include polyethylene, polypropylene, polybutylene, poly(4-methylbutylene), butyl rubber, silastic polymers, polyesters, polyamides, cellulose and cellulose derivatives (such as cellulose acetate, nitrocellulose and the like), acrylates, methacrylates, vinyl polymers (such as polyvinyl acetate, polyvinyl chloride, polyvinylidene chloride, polyvinyl fluoride, and the like), polystyrene and styrene graft copolymers, rayon, nylon, polyvinylbutyrate, polyformaldehyde, etc. Other materials which are listed are silicon wafers, glasses, insoluble protein coatings on a smooth insoluble surface, metals, metalloids, metal oxides, magnetic materials, materials used in semiconductor devices, cermets and the like. The supports disclosed as preferred include polished single crystalline silicon, aluminum, epitaxial silicon coatings, silicon nitride coatings, silicon dioxide coatings, and polysilicon coatings.

This invention is directed to improved, reflective diffraction biogratings (diffraction assay devices) suitable for use in the apparatus and methods of U.S. Patents 4,647,544 and 4,876,208. These biogratings have a higher reflectivity, a high binding capacity and optical flatness.

In summary, the biograting consists of an optically flat layer of silicon dioxide having a first and second surface, alternating zones of active and inactive binding reagent on the first surface forming a spatially periodic pattern, and a reflective metal layer having a thickness which is sufficient to prevent transmission of substantially all light (less than one percent) therethrough. The reflective metal layer can be supported on an optically flat surface of metal wafer, and the reflective metal can be aluminum, gold, silver, chromium, platinum, titanium or nickel coating on a polished wafer. Preferably, the silicon dioxide layer is formed by sputtering a layer of silicon dioxide or by coating an alkali metal silicate solution on the surface of the reflective metal. The sputtering can be carried out using conventional sputtering devices and processes, and the thickness of the silicon dioxide can be controlled by varying the discharge time. The alkali metal silicate solution optimally contains from 1 to

20 wt.% and preferably from 5 to 10 wt.% silicon dioxide; from 0.5 to 15 wt.% and preferably from 5 to 10 wt.% of an aminoalkyltrialkoxysilane; and from 1 to 20 and preferably from 5 to 10 mg/ml of a water-soluble polysaccharide. The method for making the biograting comprises uniformly adhering a binding reagent to one surface of an optically flat layer of silicon dioxide, the reflective metal layer being on the second surface; and selectively deactivating zones of the binding reagent to form spatially periodic patterns of active and deactivated binding reagent by exposing the surface to a deactivating amount of UV light through a transparent mask having a spatially periodic pattern of opaque zones thereon.

An alternative reflective biograting embodiment for a diffraction bioassay device of this invention comprises a spatially periodic pattern of zones of active and inactive binding reagent on an optically flat surface of a multilayer dielectric mirror.

Brief Description of the Drawings

Fig. 1 is a fragmentary, magnified cross-sectional view of one embodiment of this invention.

Fig. 2 is a schematic representation of the process for manufacturing an insoluble support with the spatially periodic pattern of Fig. 1.

Fig. 3 is a cross-sectional view of a dipstick having mounted thereon, a plurality of insoluble supports with spatially periodic pattern of binding reagents on the surfaces thereof.

Fig. 4 is a fragmentary, magnified cross-sectional view of an alternate embodiment of this invention with a spatially periodic pattern of binding reagent on a multilayer dielectric mirror.

Fig. 5 is a dose response curve showing data obtained in Example 11.

In an effort to increase the sensitivity of diffraction immunoassay systems using reflective biogratings, a wide variety of surfaces were investigated. The optimum reflective biograting combines high protein binding capacity, optical flatness, a high reflectivity, and selected thickness of the transparent layers. Examples of typical normal incidence reflectivities of biogratings found were bare silicon (23%), silicon oxide on silicon (12%), silicon nitride on silicon (1%), glass microscope slide (4%), and polystyrene (7%). With the improved biogratings of this invention, reflectivities greater than 60 percent have been obtained, substantially increasing the sensitivity of the bioassays developed using the improved biograting.

The term "biograting", as used herein, is defined to be a substantially flat surface having a coating thereon and having substantially uniform light scattering properties. The coating comprises a spatially periodic pattern of active and deactivated binding reagent. One suitable pattern comprises alternating, parallel linear zones of an active and deactivated binding reagent. Any other spatially periodic pattern including those described in U.S. Patent 4,647,544 can be used. The zones form a dif-

fraction grating when the active binding reagent binds with its opposite member of the binding pair. In the absence of such binding, no significant light diffraction occurs, that is, light energy detected at the diffraction angles is at a minimum value, approaching zero. When the binding occurs, the accumulation of bound material in the spatially periodic patterns create a spatially periodic light disturbing pattern, and light detected at the light diffraction angles increases to a large value which correlates to the presence and quantity of the binding partner (analyte) in the sample.

The term "binding reagent" is used herein to designate one member of any binding pair of compounds or materials which selectively bind to form a conjugate. The members of the binding pair are generically denoted by the terms "ligand" and "antiligand", either one of which can be a binding reagent. The binding reagent can be a member of the well-known antibody-antigen or antibody-hapten pairs wherein the antibody binds selectively with the respective antigen or hapten, or combinations where the antibody is replaced with an Fab, Fab', F(ab')₂ fragment or hybrid antibody. The binding reagent can also be a member of other types of binding pairs such as biotin-avidin; lectin-sugar; IgG antibody Fc portion with protein A or protein G; enzyme-enzyme substrate; DNA or RNA binding with DNA, DNA fragments or other nucleotide sequences; enzyme-enzyme inhibitor; protein-protein receptor; chelating agent-ligand; and the like. Also included are specific binding pairs wherein a mercapto group binds specifically with a dithio or disulfide group (-S-SH or -S-S-) or with a N-substituted-2,4-diketo-3-pyrroline group, and other molecules with functional groups that will bind each other specifically. In general, the binding reagent is selected to bind specifically or selectively with the analyte, the material for which a sample is assayed. A non-light disturbing layer or coating of binding reagent is applied to an insoluble surface and is transformed into a spatially periodic design of non-light disturbing material for use in the method of this invention.

The term "binding assay", is used herein to designate an assay using any binding reaction between a binding reagent and the other member of the binding pair which is selectively bindable therewith.

The term "light disturbing", as used herein, is defined to include all ways in which light is affected including light absorbing, reflecting, scattering, refracting and phase changing.

The term "spacially periodic pattern", as used herein, is defined to include gratings having spatially periodic designs which are formed in one or more immunological steps. For the method of this invention, the patterns are formed directly by the conjugation of the non-light disturbing binding reagent on the insoluble surface with a light disturbing analyte. Types of patterns formed in the process of this invention include reflection amplitude gratings, transmission amplitude gratings, reflection phase gratings, and transmission phase gratings. In re-

flexion amplitude gratings, light is reflected from the grating, and the amplitude of the reflected light is modulated by the spatially variable reflection of the grating. In transmission amplitude gratings, light is transmitted through the grating, and the amplitude of the transmitted light is modulated by the spatially variable transmission of the grating. In the reflection phase grating, the light is reflected from the grating, and the phase of the reflected light is modulated by the spatially variable refractive index of the grating. In the transmission phase gratings, light is transmitted through the grating, and the phase of the transmitted light is modulated by the spatially variable refractive index of the grating. In the method of this invention, the diffraction grating may function as one or more of these types of gratings concurrently, and all of these grating types are included within the diffraction gratings made in the method of this invention.

The term "optically flat", as used herein, is defined to be a surface with a maximum height variation of less than 600 Å over a surface area of 4 mm² or the area illuminated by the laser beam, whichever is smaller.

The term "wafer", as used herein, is defined to be a flat plate of insoluble solid having optically flat areas.

The term "alkyl", as used herein includes saturated and unsaturated, straight, branch-chained and cyclic hydrocarbon groups. The term "lower alkyl" is defined to include alkyl groups having from 1 to 6 carbon atoms.

Fig. 1 is a fragmentary, magnified cross-sectional view of one embodiment of this invention. It consists essentially of an optically flat layer of a transparent material 2 having a first surface 4 and a second surface 6. It has a spatially periodic pattern of zones of active binding reagent 8 and inactive binding reagent 10 on the first surface. It also has a reflective metal layer 12 on the second surface 6 having a thickness sufficient to prevent transmission of substantially all light (less than one percent) therethrough. For aluminum, for example, the reflective metal layer 12 should have a thickness of at least about 1000 Å. The active binding agent zones 8 have a width, a, and a distance between centers of the binding agent or "period", d.

The reflective metal layer 12 can be any reflective metal which has the stability required for the processing steps and an inherent reflectivity (for polished or optically flat surfaces) of at least 40%. Examples of suitable reflective metals include aluminum, gold, silver, chromium, titanium, nickel and platinum.

The transparent layer or coating 2 can be any transparent material which can bind protein and can be applied as a coating. It can be an organic material such as an organic polymer such as nitrocellulose. It can also be an inorganic material such as a silicon dioxide. The invention is hereinafter described with the use of silicon dioxide for purposes of clarity of description and not by way of limitation. Any transparent material satisfying the above requirements can be used and are considered to be within the scope of this invention.

The silicon dioxide layer can be any plate with op-

tically flat areas of transparent glass containing silicon dioxide, preferably treated with a suitable silane to increase its protein binding capacity. If the silicon dioxide is a self-supporting layer such as a microscope slide or coverslip, for example, the reflective metal can be applied to the surface opposite to the side carrying the biograting with a mirroring, vapor deposition, sputtering or other metallization process.

The combination yielding the highest optical flatness and reflectivity comprises a silicon dioxide coating formed on a reflective metal coated, optically flat area of wafer 14 of silicon or silicon dioxide. The wafer is the supporting layer and should have the physical and chemical stability to undergo the metallization process without significant change. A convenient source of wafers are polished plates of semiconductor materials such as silicon wafers typically used in semiconductor manufacture. These are readily available in a polished, optically flat form and have the thermal and chemical stability for metallization by vapor deposition or metal sputtering, both conventional and well known processes commonly used in semiconductor manufacture. However, polished glass would be equally suitable as a substrate since no radiation penetrates the metal layer.

The optimum reflective metal coating process depends upon the particular metal used. Aluminum, gold and silver coatings can be directly applied to one surface of the support wafer in a sputtering process carried out in an inert atmosphere, usually in a partial vacuum. The coating thickness is controlled to be sufficient to reflect all (at least 99 %) of the incident light. The thickness of reflective aluminum, for example, required to achieve this result is about 1000 Å. Suitable processes for depositing the metal coatings, for example sputtering and vapor deposition, are described in VLSI TECHNOLOGY, Edited by S.M. Sze, New York: McGraw-Hill (1983).

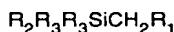
The silicon dioxide coating is then applied to the reflective metal surface by a process which yields a product having a high reflectivity and an optically flat surface.

The silicon dioxide coating can be applied by sputtering, as described in VLSI TECHNOLOGY (supra, p 358). Alternatively, the silicon dioxide coating can be applied by spin coating the reflective surface with an alkali metal silicate solution. Spin coating is a conventional process, well known to a person skilled in the coating art. The thickness of the coating is determined by the viscosity of the alkali metal silicate solution, spinning speed, temperature and evaporation rate. In general, the surface is spun around an axis perpendicular to the surface, and the solution is applied either before or during the spinning. If the coating is applied to a conventional circular wafer disk having a diameter of 4 inches, the spinning speed should be from 1500 to 8000 rpm and preferably is from 2500 to 4000 rpm.

The alkali metal silicate solution can be made of any alkali metal (sodium, potassium, lithium, etc) and is preferably a conventional sodium silicate (water glass) solution containing from 1 to 20 wt.% and preferably from

5 to 10 wt.% alkali metal silicate.

The protein binding capacity of the silicon dioxide product is increased if the surface is treated with a protein binding reagent such as an aminosilane. If the silicon dioxide coating is formed from an alkali metal silicate solution, the protein binding reagent can be incorporated directly in the coating solution. Suitable aminosilanes include aminoalkylsilanes having the formula:



wherein,

R_1 is hydrogen, an aminoalkyl group having from 1 to 18 carbons, or an aminoalkylamino group having from 1 to 18 carbons; and

R_2 , and R_3 are each, individually, a lower alkyl or alkoxy group.

Examples of suitable aminoalkylsilanes include aminopropyltriethoxysilane, aminopropyltrimethoxysilane, aminobutyltriethoxysilane, N-(2-aminoethyl-3-aminopropyl)triethoxysilane, ω -aminoundecyltrimethoxysilane, and aminopropylmethyldiethoxysilane, for example. A preferred aminoalkyltrialkoxysilane is N-(2-aminoethyl-3-aminopropyl)triethoxysilane. The alkali metal silicate solution can contain from 0.5 to 15 wt.% and preferably from 5 to 10 wt.% of the aminoalkylsilane.

The binding capacity is further increased if the alkali metal silicate solution also includes from 1 to 20 mg/ml and preferably from 5 to 10 mg/ml of a water soluble hydroxylated polymers, preferably polysaccharides. Suitable polysaccharides include water-soluble gums, hydrolyzed starches, cellulose derivatives, and other conventional water-soluble hydroxylated polymers. A particularly suitable polysaccharide are the dextrans having a molecular weight of from 5000 to 500,000 and preferably from 10,000 to 75,000.

The silicon dioxide coating thickness is determined by the sputtering time in the sputter coating process or by the speed of rotation in the spin coating process.

If the silicon dioxide coating is applied as an alkali silicate solution, the coated support is cured by heating in an oven at a temperature of from 90° to 200°C and preferably from 120° to 150°C for a time sufficient to cure the coating. The heating time will depend upon the thickness of the coating and the concentration of the coating solution. A heating time of from 0.5 to 16 hours is sufficient. A heating time of from 1 to 3 hours is preferred.

The method for making the biograting comprises a first step of uniformly adhering a binding reagent to the silicon dioxide surface. This followed by a step of selectively deactivating zones of the binding reagent to form a spatially periodic pattern of active and deactivated

binding agent by exposing the surface to a deactivating amount of UV light through a transparent mask having a diffraction grating pattern of opaque zones thereon.

The binding reagent applied to the silicon dioxide surface of an insoluble support is selected to bind with the analyte to be determined in the assay. It can be any member of the binding pairs described above. It can be an antibody; antibody fragment selected from the group consisting of Fab, Fab', or F(ab')₂ fragments; hybrid antibody; antigen; hapten; protein A; protein G; lectin; biotin; avidin; chelating agent; enzyme; enzyme inhibitor; protein receptor; nucleotide hybridizing agent; or a bacteria, virus, Mycoplasma, spore, parasite, yeast, or fragment thereof; or combinations thereof.

The combined optical thickness or optical path length of the transparent layer 2 and the binding layers (8 and 10) applied thereto is preferably about one-fourth the wavelength ($\lambda/4$) of the laser light to be used, measured in the direction of the laser light.

Fig. 2 is a schematic representation of the process for manufacturing an insoluble support with the diffraction grating design of Fig. 1. One member of the binding pair can be applied to the silicon dioxide surface 20 (Step A) by covalent bonding or adsorption in solution 22 in Step B. For covalent bonding, the surface, after being coated with an aminosilane, can be reacted with the protein.

One procedure for conjugating aminosilane groups with proteins can be achieved with 1-ethyl-3-(3-dimethylaminopropyl)-carbodiimide (EDCI). This is a water-soluble carbodiimide which is used for coupling antibodies or proteins with haptens or solid phases through functional groups such as carboxy and/or amino groups. EDCI reactions can be carried out as follows: To a solution of antigen or antibody in 0.01 M phosphate buffered saline, pH 6.0 at 4° C, is added an excess amount (normally 100 times the stoichiometric amount) of EDCI. The insoluble support having amino groups thereon, is added to the solution. After addition, the mixture is stirred at refrigerated temperatures for 16-24 hours to complete the reaction.

If the protein to be coupled to the support is an antibody, the conjugation is preferably carried out with a soluble periodate such as an alkali metal periodate. To a solution of antibody in 0.2 M acetate buffer, pH 5, is added a solution of the periodate (11.2 mg of periodate per 1 mg of antibody). The mixture is stirred at 4-8 °C for 1 to 1.5 hours. It is then dialyzed against 0.1 M carbonate buffer, pH 8.9, and the resulting solution is incubated with the insoluble support in a refrigerator overnight.

The antibodies can also be coupled to the insoluble surface through a thioether linkage. In this procedure, the aminopropyltriethoxysilane activated surface is allowed to react with an excess amount of iodoacetic anhydride or bromoacetic anhydride in anhydrous dimethylformamide at room temperature overnight while protecting the reactants from light. The activated surface is

then washed thoroughly with deionized water and kept protected from light exposure until it is reacted with the antibody. Before being reacted with the iodoacetyl or bromoacetyl activated surface, the antibody is treated with 2-aminoethanethiol at 37°C for 2 hours in a degassed 0.1 M phosphate buffer solution, pH 6.0. After the reaction with 2-aminoethanethiol, the solution is chromatographically purified with a Sephadex column (Pharmacia) to remove the excess amount of 2-aminoethanethiol. The antibody reaction product has mercapto groups. It can be coupled to the solid surface by reacting the solution with the iodoacetyl or bromoacetyl activated surfaces at refrigerated temperatures overnight.

Non-covalent bonding can be achieved by immersing the surface in an aqueous buffer solution. The buffered binding reagent solution is placed in a container containing the silicon dioxide surface and incubated at room temperature until adsorption occurs, for example for from 0.5 to 18 hours and preferably from 1 to 3 hours, at temperatures of from 4 to 40°C and preferably from 20 to 26°C. The surface is then rinsed with a buffered saline solution and dried.

The concentration of binding reagent in the buffer solution is selected to provide the desired reagent density on the silicon dioxide surface. The binding reagent solution can contain from 0.02 to 100 micrograms/ml of the binding reagent and preferably contains from 10 to 50 micrograms/ml of the binding reagent in a buffered solution having a pH of from 6.0 to 9.5 and preferably from 7.0 to 8.5. The surface with the coating 26 thereon is then rinsed and dried.

A suitable rinse solution is an aqueous phosphate buffer solution such as is described in U.S. Patent 4,528,267 having a phosphate molarity of from 0.0001 to 0.05, a pH of from 6 to 8 and containing from 0.001 to 0.1 weight percent non-ionic surfactant and from 0.0001 to 0.5 weight percent of an animal serum albumin. Suitable non-ionic surfactants include polyoxyethylene ethers (BRIJ®) such as lauryl, cetyl, oleyl, stearyl, and tridecyl polyoxyethylene ethers; polyoxyethylenesorbitans (TWEEN®) such as polyoxyethylenesorbitan monolaurate, monopalmitate, monostearate, monooleate and trioleates; and other polyoxyethylene ethers (TRITON®), for example. Preferred non-ionic surfactant are the polyoxyethylenesorbitans such as polyoxyethylenesorbitan monolaurate (TWEEN 20).

A mask is prepared by photographic methods conventional in semiconductor manufacturing. For example, a mask for a linear spatially periodic pattern can be prepared on a quartz glass or other UV-transparent plate through a photoresist process similar to photography. The linear dark zones of the mask correspond to active binding reagent areas desired on the ultimate surface.

In Step C, the mask 24 is mounted in a suitable support 25 of a UV light focusing apparatus such as a Karl Suss Model 40 Mask Aligner (Karl Suss, Waterbury

Center, Vermont 05677). The mask 24 is placed over the silicon dioxide surface 20 having a coating 26 of binding reagent, and the surface is exposed to ultraviolet radiation from UV radiation source 28 until the binding capability of the portions of the binding reagent exposed to the radiation is substantially reduced or preferably eliminated. To manufacture a precision grating design, the radiation should form a sharp image on the coated surface. Penumbrae should be minimized. Preferably, the ultraviolet light passing through the mask is focused to a sharp image on the surface coating using conventional projection alignment techniques without contact with the coated surface.

The ultraviolet radiation exposure required to deactivate coating exposed thereto depends upon the binding reagent. For antibody binding reagents, exposure times of from 30 sec to 30 min and preferably from 1 to 5 min is sufficient with a ultraviolet radiation having a wavelength such as 254 nm and a power of from 8 to 14 milliwatts per cm². Some adjustment in time of exposure and/or power may be necessary to deactivate the binding sites of other binding reagents.

To alter the epitopes of antigenic binding reagents such as human IgG, exposure times of from 5 to 30 min and preferably from 5 to 10 min are sufficient with a ultraviolet radiation having a wavelength of 254 nm and a power of from 8 to 14 milliwatts per cm². Some adjustment in time of exposure and/or power may be necessary to alter or destroy the antigenic sites of other binding reagents.

This treatment reduces or eliminates the binding properties of the binding reagent in zones 10, leaving active binding reagent in a spatially periodic pattern or design as the zones 8 (Fig. 1).

In Step D, the coated substrate containing areas having binding protein in a diffraction grating design is cut into smaller area chips 30, each chip having a size sufficient to perform a binding assay. These chips are then mounted on a suitable diagnostic support such as the dipstick shown in Fig. 3.

Fig. 3 is a cross-sectional view of a dipstick having mounted thereon, a plurality of insoluble supports with non-light disturbing spatially periodic patterns or designs of binding reagents on the surfaces thereof. The dipstick body 32 has a plurality of insoluble support surfaces 34 having a spatially periodic pattern or design of binding reagent coated thereon such the biogratings shown in Fig. 1 made by the process shown in Fig. 2. The materials from which the dipstick 32 are made are preferably non-binding to minimize non-specific binding during the binding assay procedure. Suitable dipstick surface materials include polyolefins such as polyethylene and polypropylene, hydrophilic polysilicon and polysiloxane polymers, and the like. Also suitable are polymers which have been treated to render the surfaces non-binding to proteinaceous materials. The silanes can be applied to the silicon dioxide surface in a vapor phase, for example.

The support for the diffraction grating supports can be any articles upon which the diffraction grating support surface can be mounted. The description of dipsticks are provided by way of example, and not as a limitation. Other articles such as microwells, plates, cavities and the like can be used. For many applications, dipsticks are a preferred embodiment.

Fig. 4 is a fragmentary, magnified cross-sectional view of an alternate embodiment of this invention with a spatially periodic pattern of binding reagent on a multilayer dielectric mirror. This highly reflective device comprises alternating layers of two transparent materials having different refractive indexes. Multiple dielectric mirrors and suitable transparent materials for manufacturing them are described in Born, Max et al, PRINCIPLES OF OPTICS. 6th ed. New York: Pergamon Press, pp 66 et seq (1980) and Hecht, Eugene, OPTICS. 2nd ed. Reading: Addison-Wesley, pp 377 et seq, the entire contents of which are hereby incorporated by reference. The preferred optical thickness of each layer is approximately one-quarter wavelength ($\lambda/4$) of the light to be reflected measured in the direction of the light. The mirrors are thus manufactured for use with collimated light of a selected frequency, usually a selected laser frequency.

The multilayer dielectric mirror comprises alternating layers of units of a layer of one transparent material such as silicon dioxide 70 and a layer of another transparent material such as titanium dioxide 72 having the quarter wavelength optical thicknesses described above. The bottom reflective layer unit is supported on the optically flat surface of a wafer 74. Each successive reflective layer unit provides an incremental increase to the reflectivity. Although no theoretical limit to the number of reflective layer units exists, the amount of the reflectivity increase provided with each additional unit is incrementally smaller.

As described above with respect to the embodiment of Fig. 1, the binding layer provides a spatially periodic design or pattern of active binding reagent. The periodic design comprises a plurality of zones of non-light disturbing active binding agent 76 separated by zones of non-light disturbing deactivated binding reagent 78, for example, binding reagent which has been deactivated by exposure to ultraviolet radiation, other deactivating radiation, or other deactivation energy. The binding reagent is a member of a binding pair as described above.

This invention is further illustrated by the following specific but non-limiting examples. Examples which have been reduced to practice are stated in the past tense, and examples which are constructively reduced to practice herein are presented in the present tense. Temperatures are given in degrees Centigrade and weight as weight percents unless otherwise specified.

EXAMPLE 1

Aluminum Coated Silicon Surface

5 Bare silicon wafers are placed in a high frequency magnetron sputtering chamber, the chamber evacuated to 5×10^{-6} torr, argon gas is introduced, and the plasma glow discharge is initiated to deposit aluminum on the silicon wafer surface using the procedures of VLSI TECHNOLOGY (supra) and MICROCHIP FABRICATION: A PRACTICAL GUIDE TO SEMICONDUCTOR PROCESSING, (supra). After 35 minutes, the plasma is terminated, the chamber vented, and the aluminum coated wafer removed.

EXAMPLE 2

Silicon Dioxide Coated Al/Si Support

20 The product of Example 1 is placed in a high frequency magnetron sputtering chamber, the chamber evacuated to 5×10^{-6} torr, and plasma gas is introduced. A plasma glow is initiated to clean the aluminum surface. Then a silicon dioxide target is introduced or exposed, and after the desired silicon dioxide coating is formed on the aluminum surface, about 30 minutes, the plasma is terminated, the vessel vented, and the aluminum-on-silicon wafers with a 1000 Å coating of silicon dioxide are removed.

EXAMPLE 3

APTS coated $\text{SiO}_2/\text{Al/Si}$ Support

35 Aminopropyltriethoxysilane (APTS) is coated onto a silicon dioxide/aluminum/silicon wafer by vapor deposition. The wafer is placed in a vacuum oven, the oven heated to 170°C , and the chamber evacuated to about 0.1 torr. The aminopropyltriethoxysilane is introduced into the oven, allowed to vaporize, and the coating process is continued for 4 hr. The oven is evacuated to remove remaining aminopropyltriethoxysilane vapor, and the wafer is retained in the oven for 8 hr to complete the coating reaction. The oven is then vented, and the APTS coated $\text{SiO}_2/\text{Al/Si}$ wafer is removed.

EXAMPLE 4

Sodium Silicate Coating on Al/Si Support

50 A 3.1 molar sodium silicate solution (VMR Scientific Catalog No. AL 68330) was diluted 1:4 with deionized water, and pipetted onto an aluminum coated silicon wafer. The wafer was spun with a Model 6000 spin coater (Integrated Technologies, Inc. Asushnet, MA 02743) at 3000 rpm for one min. The silicate coated wafer was then cured in an oven at 135°C for 2 hr and allowed to cool to room temperature.

EXAMPLE 5

APTS/SILICATE coating on Al/Si Support

A mixture of 3.1 molar sodium silicate solution, aminopropyltriethoxysilane, and deionized water (1:0.5:3.5 v/v/v) was prepared. The solution was pipetted onto an aluminum coated silicon wafer. The wafer was spun with a Model 6000 spin coater (Integrated Technologies, Inc. Asushnet, MA 02743) at 3000 rpm for one min. The silicate coated wafer was then cured in an oven at 135°C for 2 hr and allowed to cool to room temperature.

EXAMPLE 6

APTS-DEXTRAN-SILICATE Coating on Al/Si Support

To each ml of a mixture of 3.1 molar sodium silicate solution, aminopropyltriethoxysilane, and deionized water (1:0.5:3.5 v/v/v) was added 5 mg of dextran (50,000 daltons). The mixture was pipetted onto a aluminum coated silicon wafer. The wafer was spun with a Model 6000 spin coater (Integrated Technologies, Inc. Asushnet, MA 02743) at 3000 rpm for one min. The silicate coated wafer was then cured in an oven at 135°C for 2 hr and allowed to cool to room temperature.

EXAMPLE 7

Reflectivities of Coated Al/Si Supports

The reflectivities of the silicon surfaces prepared in Examples 1-6 were determined. The silicon surfaces were illuminated with a Model 1107P Helium-Neon Laser (Uniphase, Sunnyvale, CA 94086) at an incident angle of 75°. The reflected light intensity was measured with a Model 61 Optometer (United Detector Technology, Hawthorne, CA). A reflectivity of 80% or higher was found on all surfaces.

Surface Coating	Reflectivity
Silicon Dioxide	0.92
APTS/Silicon Dioxide	0.84
Silicate	0.85
APTS/Silicate	0.80
APTS/Silicate/Dextran	0.80

EXAMPLE 8

Monoclonal Anti-β-hCG (Fab) coating

A silicon dioxide/aluminum/silicon wafer, coated by vapor deposition with APTS, was incubated with a solution of monoclonal anti-β-hCG (Fab) in 0.01 M phosphate buffer, pH 7.4 (100 μg/ml) at 4-8°C for 4 hr. The surface was briefly washed with 0.05 M Tris buffer, pH

8.5, containing 2.5% sucrose. It was then incubated with 0.05 M Tris buffer, pH 8.5, containing 2.5% sucrose and 0.5 wt.% human serum albumin (HSA) at 4-8°C for 30 min. The residual liquid was removed by spinning the wafer to yield an anti-β-hCG coated silicon dioxide surface.

EXAMPLE 9

Monoclonal Anti-β-hCG (IgG) Coating

Monoclonal anti-β-hCG antibody was diluted in 0.2 M of acetate buffer, pH 5.0, to a concentration of 2 mg/ml. The solution was cooled in an ice bath, and a solution of sodium periodate (22.4 mg/ml) in 0.1 M acetate buffer was slowly added. The mixture was then stirred at 4-8°C for 1 to 1.5 hr and then dialyzed against 2 L of 0.1 M carbonate buffer, pH 8.9, at 4-8°C for 4-6 hr. The antibody solution obtained was incubated with a silicon dioxide/aluminum coated silicon wafer surface which has been coated with APTS by vapor deposition in a refrigerator overnight. The surface was briefly washed with 0.05 M Tris buffer, pH 8.5, containing 2.5% sucrose. It was then incubated with 0.05 M Tris buffer, pH 8.5, containing 2.5% sucrose and 0.5 wt.% human serum albumin (HSA) at 4-8°C for 30 min. The residual liquid was removed by spinning the wafer to yield an anti-β-hCG coated silicon dioxide surface.

EXAMPLE 10

Biograting Preparation

An anti-β-hCG coated silicon dioxide surface prepared by the procedure of Example 8 was placed under a photomask using a Karl Suss Model 40 Mask Aligner. The photomask has parallel opaque lines having a center-to-center distance, d, of 10 μm. The surface was illuminated with UV light at 254 nm for 6 min. After illumination, the surface was diced into 4 x 6 mm chips.

EXAMPLE 11

Anti-β-hCG (Fab) Biograting Immunoassay

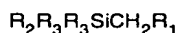
The 4 x 6 mm chips prepared in Example 10 were mounted on a microscope slide and incubated with 600, 300, 150, 75 and 0 mIU/ml of β-hCG solutions (test samples) at room temperature for 5 min. The surface was washed with deionized water and dried with a stream of air. The diffraction intensities of the surfaces were determined with a Model 61 optometer to give a dose response curve shown in Fig. 5.

Claims

1. A reflective diffraction assay device for detecting an

analyte comprising an optically flat layer of a transparent composition having a first and second surface, a spatially periodic pattern of zones of active and inactive binding reagent on the first surface, and a reflective metal layer on the second surface, the reflective metal layer having a thickness sufficient to prevent transmission of substantially all incident light therethrough, wherein said active binding reagent selectively binds said analyte.

2. The reflective diffraction assay device of Claim 1 wherein the reflective metal layer is supported on an optically flat surface.
3. The reflective diffraction assay device of Claim 1 or Claim 2 wherein the reflective metal is aluminum, gold, silver, chromium, platinum, nickel or titanium.
4. The reflective diffraction assay device of any one of the preceding claims wherein the transparent composition comprises silicon dioxide.
5. The reflective diffraction assay device of Claim 4 wherein the layer of silicon dioxide is formed by sputtering of silicon dioxide.
6. The reflective diffraction assay device of any one of claims 1 to 3 wherein transparent composition is formed by coating an alkali metal silicate solution on the surface of the reflective metal.
7. The reflective diffraction assay device of claim 4 or claim 5 wherein the first surface of the transparent layer comprises a protein binding reagent.
8. The reflective diffraction assay device of claim 7 wherein the transparent composition is formed by coating an alkali metal silicate solution on the surface of the reflective metal, and the protein binding reagent is present in the solution.
9. The reflective diffraction assay device of claim 7 or claim 8 wherein the protein binding reagent is an aminosilane.
10. The reflective diffraction assay device of claim 9 wherein the aminosilane is an aminoalkylsilane.
11. The reflective diffraction assay device of claim 10 wherein the aminosilane is an aminoalkylsilane having the formula:



wherein,

R_1 is hydrogen, an aminoalkyl group having

from 1 to 18 carbons, or an aminoalkylamino group having from 1 to 18 carbons; and

R_2 , and R_3 are each, individually, a lower alkyl or alkoxy group.

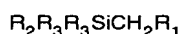
12. The reflective diffraction assay device of Claim 11 wherein the transparent composition layer is formed by coating an alkali metal silicate solution on the surface of the reflective metal, and the alkali metal silicate solution contains from 1 to 20 wt.% alkali metal silicate and from 0.5 to 15 wt.% of the aminoalkylsilane.
13. The reflective diffraction assay device of Claim 11 wherein the aminoalkylsilane is applied and bonded to the layer of silicon dioxide by vapor deposition.
14. The reflective diffraction assay device of any one of claims 11 to 13 wherein the aminoalkylsilane is aminopropyltriethoxysilane or N-(2-aminoethyl)-3-aminopropyltriethoxysilane.
15. The reflective diffraction assay device of Claim 12 wherein the alkali metal silicate solution contains from 1 to 20 mg/ml of a water-soluble hydroxylated polymer.
16. The reflective diffraction assay device of Claim 15 wherein the hydroxylated polymer is a dextran having a molecular weight in the range of from 5000 to 500,000.
17. The reflective diffraction assay device of any one of the preceding claims wherein the binding reagent is a member of a binding pair, one member of which is an antibody; antibody fragment selected from the group consisting of Fab', Fab, F(ab')₂ fragments; hybrid antibody; antigen; hapten; protein A; protein G; lectin; biotin; avidin; chelating agent; enzyme; enzyme inhibitor; protein receptor; nucleotide hybridizing agent; or a bacterium, virus, Mycoplasma, spore, parasite, yeast, or fragment thereof; or combinations thereof.
18. A method for making a reflective diffraction assay device comprising

a) uniformly adhering a binding reagent to a first surface of an optically flat layer of a transparent composition having a first and second surface, a reflective metal layer on the second surface having a thickness sufficient to prevent transmission of substantially all light therethrough; and

b) selectively deactivating zones of the binding reagent to form a diffraction grating pattern of

alternating zones of active and deactivated binding reagent by exposing the surface to a deactivating amount of UV light through a transparent mask having a spatially periodic pattern of opaque zones thereon.

19. The method of Claim 18 wherein the reflective metal layer is supported on an optically flat surface.
20. The method of Claim 18 or Claim 19 wherein the reflective metal is aluminium, gold, silver, chromium, platinum, nickel or titanium.
21. The method of any one of Claims 18 to 20 wherein the transparent composition is silicon dioxide.
22. The method of Claim 21 wherein the layer of silicon dioxide is formed by sputtering of silicon dioxide.
23. The method of any one of claims 18 to 20 wherein the transparent composition is formed by coating an alkali metal silicate solution on the surface of the reflective metal.
24. The method of Claim 21 or Claim 22 wherein the first surface of the transparent composition layer comprises an aminoalkylsilane having the formula:



wherein,

R_1 is hydrogen, an aminoalkyl group having from 1 to 18 carbons, or an aminoalkylamino group having from 1 to 18 carbons; and R_2 , and R_3 are each, individually, a lower alkyl or alkoxy group.

25. The method of claim 24 wherein the transparent composition layer is formed by coating an alkali metal silicate solution on the surface of the reflective metal, and the alkali metal silicate solution contains from 1 to 20 wt.% alkali silicate and from 0.5 to 15 wt.% of the aminoalkylsilane.
26. The method of claim 24 wherein the aminoalkylsilane is applied and bonded to the layer of silicon dioxide by vapor deposition.
27. The method of any one of claims 24 to 26 wherein the aminoalkylsilane is aminopropyltriethoxysilane or N-(2-aminoethyl)-3-aminopropyl-triethoxysilane.
28. The method of Claim 25 wherein the alkali metal silicate solution contains from 1 to 20 mg/ml of a water-soluble hydroxylated polymer.

29. The method of claim 28 wherein the hydroxylated polymer is a dextran having a molecular weight in the range of from 5000 to 500,000.

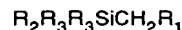
- 5 30. The method of any one of claims 18 to 29 wherein the binding reagent is a member of a binding pair, one member of which is an antibody; antibody fragment selected from the group consisting of Fab', Fab, F(ab')₂ fragments; hybrid antibody; antigen; hapten; protein A; protein G; lectin; biotin; avidin; chelating agent; enzyme; enzyme inhibitor; protein receptor; nucleotide hybridizing agent; or a bacteria, virus, Mycoplasma, spore, parasite, yeast, or fragment thereof; or combinations thereof.
- 10
- 15 31. A reflective diffraction assay device comprising alternating zones of active and inactive binding reagent in a spatially periodic pattern on the surface layer of a multilayer dielectric mirror comprising alternating layers of first and second transparent materials supported on an optically flat support, each of the first and second transparent materials having differing diffraction indexes.
- 20
- 25 32. The reflective diffraction assay device of Claim 31 wherein said surface layer comprises the first transparent material and is silicon dioxide.
- 30 33. The reflective diffraction assay device of Claim 31 or Claim 32 wherein the second transparent material is titanium dioxide.
- 35 34. The reflective diffraction assay device of any one of claims 31 to 33 wherein the binding reagent is a member of a binding pair, one member of which is an antibody; antibody fragment selected from the group consisting of Fab', Fab, F(ab')₂ fragments; hybrid antibody; antigen; hapten; protein A; protein G; lectin; biotin; avidin; chelating agent; enzyme; enzyme inhibitor; protein receptor; nucleotide hybridizing agent; or a bacterium, virus, Mycoplasma, spore, parasite, yeast, or fragment thereof; or combinations thereof.
- 40
- 45 35. The reflective diffraction assay device of any one of claims 1 to 17 wherein the reflective metal layer has a thickness sufficient to reflect at least 99% of incident light.
- 50 36. The reflective diffraction assay device of Claim 35 wherein the reflective metal layer is aluminium of a thickness of at least 1000Å.
- 55 37. The reflective diffraction assay device of any one of claims 1 to 17 wherein the transparent composition layer has a thickness of about 1000Å.

Patentansprüche

1. Reflexions-Diffraktionsassay-Vorrichtung zum Nachweis eines Analyten, umfassend eine optisch plane Schicht aus einer transparenten Zusammensetzung mit einer ersten und einer zweiten Oberfläche, ein räumlich periodisches Zonenmuster von aktivem und inaktivem Bindereagens auf der ersten Oberfläche und eine reflektierende Metallschicht auf der zweiten Oberfläche, wobei die reflektierende Metallschicht eine Dicke aufweist, die ausreicht, um den Durchgang im wesentlichen allen auftretenden Lichts durch sie hindurch zu verhindern, wobei das aktive Bindereagens den Analyten selektiv bindet.
2. Reflexions-Diffraktionsassay-Vorrichtung nach Anspruch 1, worin die reflektierende Metallschicht auf einer optisch planen Oberfläche getragen wird.
3. Reflexions-Diffraktionsassay-Vorrichtung nach Anspruch 1 oder 2, worin das reflektierende Metall Aluminium, Gold, Silber, Chrom, Platin, Nickel oder Titan ist.
4. Reflexions-Diffraktionsassay-Vorrichtung nach einem der vorangegangenen Ansprüche, worin die transparente Zusammensetzung Siliziumdioxid umfaßt.
5. Reflexions-Diffraktionsassay-Vorrichtung nach Anspruch 4, worin die Schicht aus Siliziumdioxid durch Sputtern von Siliziumdioxid gebildet wird.
6. Reflexions-Diffraktionsassay-Vorrichtung nach einem der Ansprüche 1 bis 3, worin die transparente Zusammensetzung durch Beschichten der Oberfläche des reflektierenden Metalls mit einer Alkalimetallsilikat-Lösung gebildet wird.
7. Reflexions-Diffraktionsassay-Vorrichtung nach Anspruch 4 oder 5, worin die erste Oberfläche der transparenten Schicht ein Proteinbindereagens umfaßt.
8. Reflexions-Diffraktionsassay-Vorrichtung nach Anspruch 7, worin die transparente Zusammensetzung durch Beschichten der Oberfläche des reflektierenden Metalls mit einer Alkalimetallsilikat-Lösung gebildet wird und das Proteinbindereagens in der Lösung vorhanden ist.
9. Reflexions-Diffraktionsassay-Vorrichtung nach Anspruch 7 oder 8, worin das Proteinbindereagens ein Aminosilan ist.
10. Reflexions-Diffraktionsassay-Vorrichtung nach Anspruch 9, worin das Aminosilan ein Aminoalkylsilan

ist.

11. Reflexions-Diffraktionsassay-Vorrichtung nach Anspruch 10, worin das Aminosilan ein Aminoalkylsilan der Formel



ist, worin

R_1 Wasserstoff, eine Aminoalkylgruppe mit 1 bis 18 Kohlenstoffen oder eine Aminoalkylaminogruppe mit 1 bis 18 Kohlenstoffatomen ist; und
 R_2 und R_3 unabhängig voneinander jeweils eine Niederalkyl- oder -alkoxygruppe sind.

12. Reflexions-Diffraktionsassay-Vorrichtung nach Anspruch 11, worin die Schicht der transparenten Zusammensetzung durch Beschichten der Oberfläche des reflektierenden Metalls mit einer Alkalimetallsilikat-Lösung gebildet wird und die Alkalimetallsilikat-Lösung 1 bis 20 Gew.-% Alkalimetallsilikat und 0,5 bis 15 Gew.-% des Aminoalkylsilans enthält.
13. Reflexions-Diffraktionsassay-Vorrichtung nach Anspruch 11, worin das Aminoalkylsilan durch Dampfabscheidung auf die Siliziumdioxidschicht aufgebracht und mit dieser verbunden wird.
14. Reflexions-Diffraktionsassay-Vorrichtung nach einem der Ansprüche 11 bis 13, worin das Aminoalkylsilan Aminopropyltriethoxysilan oder N-(2-Aminoethyl)-3-aminopropyltriethoxysilan ist.
15. Reflexions-Diffraktionsassay-Vorrichtung nach Anspruch 12, worin die Alkalimetallsilikat-Lösung 1 bis 20 mg/ml eines wasserlöslichen, Hydroxylgruppen aufweisenden Polymers enthält.
16. Reflexions-Diffraktionsassay-Vorrichtung nach Anspruch 15, worin das Hydroxylgruppen aufweisende Polymer ein Dextran mit einem Molekulargewicht im Bereich von 5.000 bis 500.000 ist.
17. Reflexions-Diffraktionsassay-Vorrichtung nach einem der vorangegangenen Ansprüche, worin das Bindereagens ein Teil eines Bindungspaares ist, von dem ein Teil ein Antikörper; Antikörperfragment, ausgewählt aus der aus Fab'-, Fab-, F(ab')₂- Fragmenten bestehenden Gruppe; Hybridantikörper; Antigen; Hapten; Protein A; Protein G; Lectin; Biotin; Avidin; ein Chelatbildner; Enzym; Enzyminhibitor; Proteinrezeptor; Nukleotidhybridisierungsmittel; oder ein Bakterium, ein Virus, ein Vertreter aus der Ordnung Mycoplasmatales, eine Spore, ein Parasit, eine Hefe, oder ein Fragment davon; oder ei-

ne Kombination davon ist.

18. Verfahren zur Herstellung einer Reflexions-Diffraktionsassay-Vorrichtung, umfassend:

a) das gleichmäßige Ankleben eines Bindereagens an eine erste Oberfläche einer optisch planen Schicht aus einer transparenten Zusammensetzung mit einer ersten und einer zweiten Oberfläche, wobei eine reflektierende Metallschicht auf der zweiten Oberfläche eine Dicke aufweist, die ausreicht, um den Durchgang im wesentlichen allen Lichts durch sie hindurch zu verhindern; und
b) das selektive Deaktivieren von Zonen des Bindereagens, um ein Diffraktionsgittermuster alternierender Zonen von aktivem und deaktiviertem Bindereagens zu bilden, indem die Oberfläche durch eine transparente Maske mit einem darauf ausgebildeten, räumlich periodischen Muster lichtundurchlässiger Zonen hindurch einer deaktivierenden Menge an UV-Licht ausgesetzt wird.

19. Verfahren nach Anspruch 18, worin die reflektierende Metallschicht auf einer optisch planen Oberfläche getragen wird.

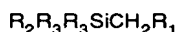
20. Verfahren nach Anspruch 18 oder 19, worin das reflektierende Metall Aluminium, Gold, Silber, Chrom, Platin, Nickel oder Titan ist.

21. Verfahren nach einem der Ansprüche 18 bis 20, worin die transparente Zusammensetzung Siliziumdioxid umfaßt.

22. Verfahren nach Anspruch 21, worin die Schicht aus Siliziumdioxid durch Sputtern von Siliziumdioxid gebildet wird.

23. Verfahren nach einem der Ansprüche 18 bis 20, worin die transparente Zusammensetzung durch Beschichten der Oberfläche des reflektierenden Metalls mit einer Alkalimetallsilikat-Lösung gebildet wird.

24. Verfahren nach Anspruch 21 oder 22, worin die erste Oberfläche der transparenten Zusammensetzung ein Aminoalkylsilan der Formel



umfaßt, worin

R_1 Wasserstoff, eine Aminoalkylgruppe mit 1 bis 18 Kohlenstoffen oder eine Aminoalkylaminogruppe mit 1 bis 18 Kohlenstoffatomen ist;

und

R_2 und R_3 unabhängig voneinander jeweils eine Niederalkyl- oder -alkoxygruppe sind.

25. Verfahren nach Anspruch 24, worin die Schicht der transparenten Zusammensetzung durch Beschichten der Oberfläche des reflektierenden Metalls mit einer Alkalimetallsilikat-Lösung gebildet wird und die Alkalimetallsilikat-Lösung 1 bis 20 Gew.-% Alkalimetallsilikat und 0,5 bis 15 Gew.-% des Aminoalkylsilans enthält.

26. Verfahren nach Anspruch 24, worin das Aminoalkylsilan durch Dampfabscheidung auf die Siliziumdioxidschicht aufgebracht und mit dieser verbunden wird.

27. Verfahren nach einem der Ansprüche 24 bis 26, worin das Aminoalkylsilan Aminopropyltriethoxysilan oder N-(2-Aminoethyl)-3-aminopropyltriethoxysilan ist.

28. Verfahren nach Anspruch 25, worin die Alkalimetallsilikat-Lösung 1 bis 20 mg/ml eines wasserlöslichen, Hydroxylgruppen aufweisenden Polymers enthält.

29. Verfahren nach Anspruch 28, worin das Hydroxylgruppen aufweisende Polymer ein Dextran mit einem Molekulargewicht im Bereich von 5.000 bis 500.000 ist.

30. Verfahren nach einem der Ansprüche 18 bis 29, worin das Bindereagens ein Teil eines Bindungspaares ist, von dem ein Teil ein Antikörper; Antikörperfragment, ausgewählt aus der aus Fab'-, Fab-, F(ab')₂-Fragmenten bestehenden Gruppe; Hybridantikörper; Antigen; Hapten; Protein A; Protein G; Lectin; Biotin; Avidin; ein Chelatbildner; Enzym; Enzyminhibitor; Proteinrezeptor; Nukleotidhybridisierungsmittel; oder ein Bakterium, ein Virus, ein Vertreter aus der Ordnung Mycoplasmales, eine Spore, ein Parasit, eine Hefe, oder ein Fragment davon; oder eine Kombination davon ist.

31. Reflexions-Diffraktionsassay-Vorrichtung, umfassend alternierende Zonen von aktivem und inaktivem Bindereagens in einem räumlich periodischen Muster auf der Oberflächenschicht eines aus mehreren Schichten bestehenden dielektrischen Spiegels, umfassend alternierende Schichten eines ersten und eines zweiten transparenten Materials, die auf einem optisch planen Träger getragen werden, wobei das erste und das zweite transparente Material jeweils unterschiedliche Brechungsindizes aufweisen.

32. Reflexions-Diffraktionsassay-Vorrichtung nach An-

spruch 31, worin die Oberflächenschicht das erste transparente Material umfaßt und aus Siliziumdioxid besteht.

33. Reflexions-Diffraktionsassay-Vorrichtung nach Anspruch 31 oder 32, worin das zweite transparente Material Titandioxid ist.

34. Reflexions-Diffraktionsassay-Vorrichtung nach einem der Ansprüche 31 bis 33, worin das Bindereagens ein Teil eines Bindungspaares ist, von dem ein Teil ein Antikörper, Antikörperfragment, ausgewählt aus der aus Fab'-, Fab-, F(ab')₂-Fragmenten bestehenden Gruppe; Hybridantikörper; Antigen; Hapten; Protein A; Protein G; Lectin; Biotin; Avidin; ein Chelatbildner; Enzym; Enzyminhibitor; Proteinrezeptor; Nukleotidhybridisierungsmittel; oder ein Bakterium, ein Virus, ein Vertreter aus der Ordnung Mycoplasmatales, eine Spore, ein Parasit, eine Heife, oder ein Fragment davon; oder eine Kombination davon ist.

35. Reflexions-Diffraktionsassay-Vorrichtung nach einem der Ansprüche 1 bis 17, worin die reflektierende Metallschicht eine Dicke aufweist, die ausreicht, um zumindest 99% des auftreffenden Lichts zu reflektieren.

36. Reflexions-Diffraktionsassay-Vorrichtung nach Anspruch 35, worin die reflektierende Metallschicht aus Aluminium mit einer Dicke von zumindest 1000 Å besteht.

37. Reflexions-Diffraktionsassay-Vorrichtung nach einem der Ansprüche 1 bis 17, worin die transparente Zusammensetzungsschicht eine Dicke von etwa 1000 Å aufweist.

Revendications

1. Dispositif d'analyse de diffraction réfléchissant pour détecter un analyte comprenant une couche optiquement plane d'une composition transparente ayant une première et une seconde surface, un motif spatialement périodique de zones d'un réactif de liaison actif et inactif sur la première surface, et une couche de métal réfléchissant sur la seconde surface, la couche de métal réfléchissant ayant une épaisseur suffisante pour empêcher la transmission de substantiellement toute la lumière incidente à travers elle, où ledit réactif de liaison actif se lie sélectivement audit analyte.

2. Dispositif d'analyse de diffraction réfléchissant selon la revendication 1 où la couche de métal réfléchissant est supportée sur une surface optiquement plane.

3. Dispositif d'analyse de diffraction réfléchissant selon la revendication 1 ou la revendication 2 où le métal réfléchissant est de l'aluminium, de l'or, de l'argent, du chrome, du platine, du nickel ou du titane.

4. Dispositif d'analyse de diffraction réfléchissant selon l'une quelconque des revendications précédentes où la composition transparente comprend du dioxyde de silicium.

5. Dispositif d'analyse de diffraction réfléchissant selon la revendication 4 où la couche de dioxyde de silicium est formée par vaporisation de dioxyde de silicium.

6. Dispositif d'analyse de diffraction réfléchissant selon l'une quelconque des revendications 1 à 3 où la composition transparente est formée par revêtement d'une solution de silicate d'un métal alcali sur la surface du métal réfléchissant.

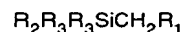
7. Dispositif d'analyse de diffraction réfléchissant selon la revendication 4 ou la revendication 5 où la première surface de la couche transparente comprend un réactif de liaison de protéines.

8. Dispositif d'analyse de diffraction réfléchissant selon la revendication 7 où la composition transparente est formée par revêtement d'une solution de silicate de métal alcali sur la surface du métal réfléchissant, et le réactif de liaison de protéines est présent dans la solution.

9. Dispositif d'analyse de diffraction réfléchissant selon la revendication 7 ou la revendication 8 où le réactif de liaison de protéines est un aminosilane.

10. Dispositif d'analyse de diffraction réfléchissant selon la revendication 9 où l'aminosilane est un aminoalkylsilane.

11. Dispositif d'analyse de diffraction réfléchissant selon la revendication 10 où l'aminosilane est un aminoalkylsilane ayant la formule :



où

R₁ est un hydrogène, un groupe aminoalkyle ayant de 1 à 18 carbones, ou un groupe aminoalkylamino ayant de 1 à 18 carbones ; et R₂, et R₃ sont chacun, individuellement, un groupe alcoxy ou alkyle inférieur.

12. Dispositif d'analyse de diffraction réfléchissant se-

lon la revendication 11 où la couche de composition transparente est formée par revêtement d'une solution de silicate de métal alcali sur la surface du métal réfléchissant, et la solution de silicate de métal alcali contient de 1 à 20% en poids de silicate de

13. Dispositif d'analyse de diffraction réfléchissant selon la revendication 11 où l'aminosilane est appliqué et lié à la couche de dioxyde de silicium par dépôt en phase vapeur.

14. Dispositif d'analyse de diffraction réfléchissant selon l'une quelconque des revendications 11 à 13 où l'aminosilane est l'aminopropyltriéthoxysilane ou le N-(2-aminoéthyl)-3-aminopropyltriéthoxysilane.

15. Dispositif d'analyse de diffraction réfléchissant selon la revendication 12 où la solution de silicate de métal alcali contient de 1 à 20 mg/ml d'un polymère hydroxylé soluble dans l'eau.

16. Dispositif d'analyse de diffraction réfléchissant selon la revendication 15 où le polymère hydroxylé est un dextran ayant un poids moléculaire dans l'intervalle de 5 000 à 500 000.

17. Dispositif d'analyse de diffraction réfléchissant selon l'une quelconque des revendications précédentes où le réactif de liaison est un membre d'une paire de liaisons, dont un membre est un anticorps ; un fragment d'anticorps choisi dans le groupe consistant en les fragments Fab', Fab, F(ab')₂ ; un anticorps hybride ; un antigène ; un haptène ; une protéine A ; une protéine G ; la lectine ; la biotine ; l'avidine ; un agent chélatant ; un enzyme ; un inhibiteur d'enzyme ; un récepteur de protéine ; un agent d'hybridation de nucléotide ; ou une bactérie, un virus, des Mycoplasmales, un spore, un parasite, une levure, ou un fragment de ceux-ci ; ou des combinaisons de ceux-ci.

18. Méthode de fabrication d'un dispositif d'analyse de diffraction réfléchissant comprenant :

a) l'adhésion uniformément d'un réactif de liaison sur une première surface d'une couche optiquement plane d'une composition transparente ayant une première et une seconde surface, d'une couche de métal réfléchissant sur la seconde surface ayant une épaisseur suffisante pour empêcher la transmission de substantiellement toute la lumière à travers elle ; et

b) la désactivation sélectivement de zones du réactif de liaison pour former un motif de réseau

de diffraction de zones alternées de réactif de liaison désactivé et actif par exposition de la surface à une quantité désactivante de lumière UV à travers un masque transparent ayant un motif spatialement périodique de zones opaques sur lui.

19. Méthode selon la revendication 18 où la couche de métal réfléchissant est supportée sur une surface optiquement plane.

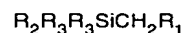
20. Méthode selon la revendication 18 ou la revendication 19 où le métal réfléchissant est l'aluminium, l'or, l'argent, le chrome, le platine, le nickel ou le titane.

21. Méthode selon l'une quelconque des revendications 18 à 20 où la composition transparente est le dioxyde de silicium.

22. Méthode selon la revendication 21 où la couche de dioxyde de silicium est formée par vaporisation de dioxyde de silicium.

23. Méthode selon l'une quelconque des revendications 18 à 20 où la composition transparente est formée par revêtement d'une solution de silicate de métal alcali sur la surface du métal réfléchissant.

24. Méthode selon la revendication 21 ou la revendication 22 où la première surface de la couche de composition transparente comprend un aminosilane ayant la formule :



où,

R₁ est un hydrogène, un groupe aminoalkyl ayant de 1 à 18 carbones, ou un groupe alkylamino ayant de 1 à 18 carbones ; et R₂, et R₃ sont chacun, individuellement, un groupe alcoxy ou alkyle inférieur.

25. Méthode selon la revendication 24 où la couche de composition transparente est formée par revêtement d'une solution de silicate de métal alcali sur la surface du métal réfléchissant, et la solution de silicate de métal alcali contient de 1 à 20% en poids de silicate alcali et de 0,5 à 15% en poids de l'aminosilane.

26. Méthode selon la revendication 24 où l'aminosilane est appliqué et lié à la couche de dioxyde de silicium par dépôt en phase vapeur.

27. Méthode selon l'une quelconque des revendications 24 à 26 où l'aminosilane et l'aminopropyl-

triéthoxysilane ou le N-(2-aminoéthyl)-3-aminopropyl-triéthoxysilane.

28. Méthode selon la revendication 25 où la solution de silicate de métal alcali contient de 1 à 20 mg/ml d'un polymère hydroxylé soluble dans l'eau.

29. Méthode selon la revendication 28 où le polymère hydroxylé est un dextran ayant un poids moléculaire dans l'intervalle de 5 000 à 500 000.

30. Méthode selon l'une quelconque des revendications 18 à 29 où le réactif de liaison est un membre d'une paire de liaisons, dont un membre est un anticorps ; un fragment d'anticorps choisi dans le groupe consistant en les fragments Fab', Fab, F(ab')₂ ; un anticorps hybride ; un antigène ; un haptène ; une protéine A ; une protéine G ; la lectine ; la biotine ; l'avidine ; un agent chélatant ; un enzyme ; un inhibiteur d'enzyme ; un récepteur de protéine ; un agent hybridant un nucléotide ; ou une bactérie, un virus, un Mycoplasmales, un spore, un parasite, une levure, ou un fragment de ceux-ci ; ou des combinaisons de ceux-ci.

31. Dispositif d'analyse de diffraction réfléchissant comprenant des zones alternées d'un réactif de liaison actif et inactif dans un motif spatialement périodique sur la couche de surface d'un miroir diélectrique multicouche comprenant des couches alternées de premier et second matériaux transparents supportés sur un support optiquement plan, chacun des premier et second matériaux transparents ayant des indices de diffraction différents.

32. Dispositif d'analyse de diffraction réfléchissant selon la revendication 31 où ladite couche de surface comprend le premier matériau transparent et est du dioxyde de silicium.

33. Dispositif d'analyse de diffraction réfléchissant selon la revendication 31 ou la revendication 32 où le second matériau transparent est du dioxyde de titane.

34. Dispositif d'analyse de diffraction réfléchissant selon l'une quelconque des revendications 31 à 33 où le réactif de liaison est un membre d'une paire de liaisons, dont un membre est un anticorps ; un fragment d'anticorps choisi dans le groupe consistant en les fragments Fab', Fab, F(ab')₂ ; un anticorps hybride ; un antigène ; un haptène ; une protéine A ; une protéine G ; la lectine ; la biotine ; l'avidine ; un agent chélatant ; un enzyme ; un inhibiteur d'enzyme ; un récepteur de protéine ; un agent hybridant un nucléotide ; ou une bactérie, un virus, un Mycoplasmales, un spore, un parasite, une levure ou un fragment de ceux-ci ; ou des combinaisons

de ceux-ci.

35. Dispositif d'analyse de diffraction réfléchissant selon l'une quelconque des revendications 1 à 17 où la couche de métal réfléchissant a une épaisseur suffisante pour réfléchir au moins 99% de la lumière incidente.

36. Dispositif d'analyse de diffraction réfléchissant selon la revendication 35 où la couche de métal réfléchissant est de l'aluminium d'une épaisseur d'au moins 1000 Å.

37. Dispositif d'analyse de diffraction réfléchissant selon l'une quelconque des revendications 1 à 17 où la couche de composition transparente a une épaisseur d'environ 1000 Å.

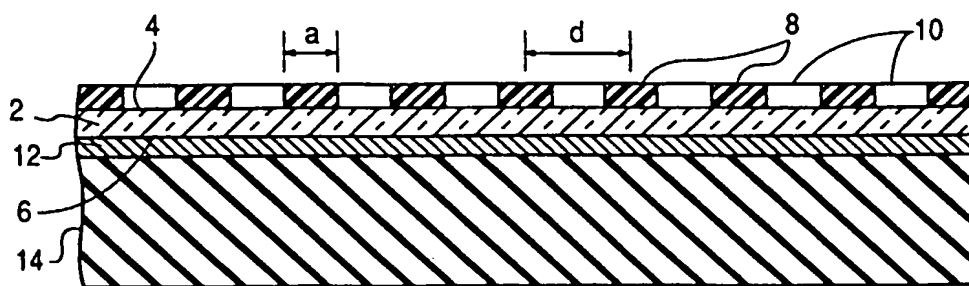


FIG. 1

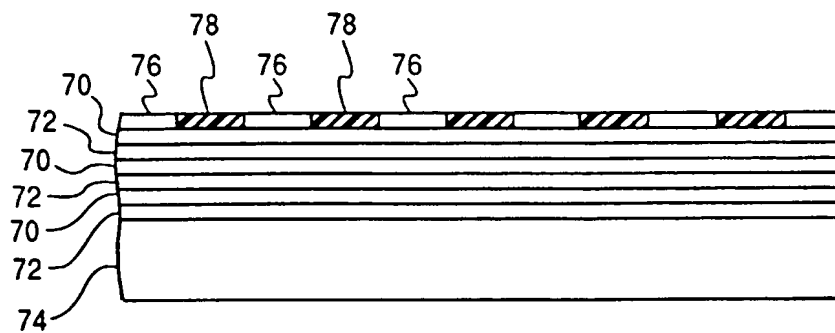


FIG. 4

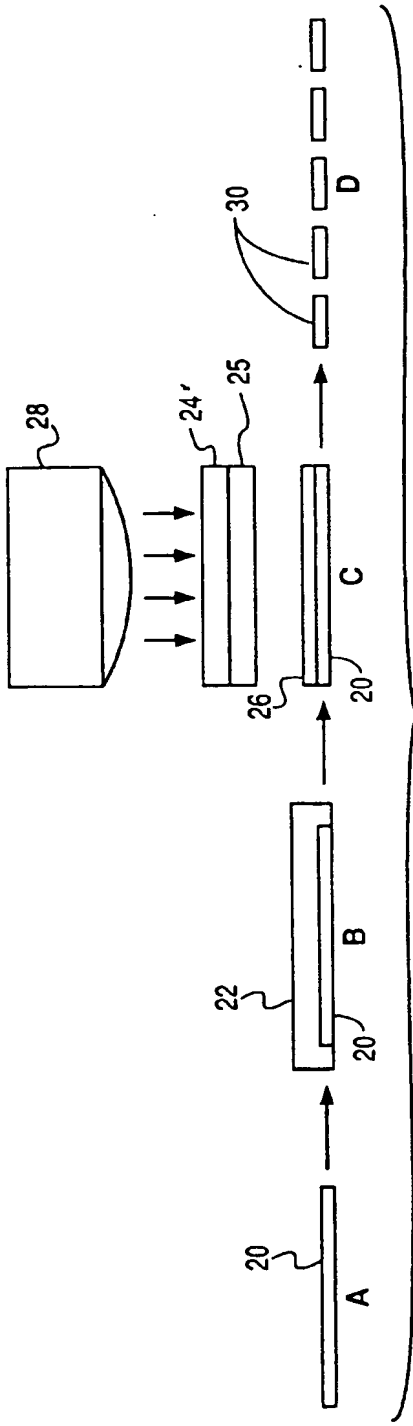


FIG. 2

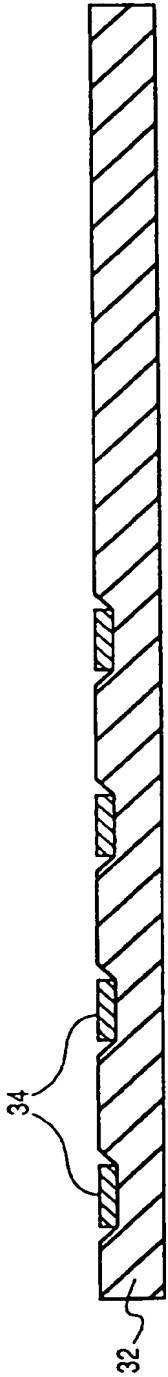


FIG. 3

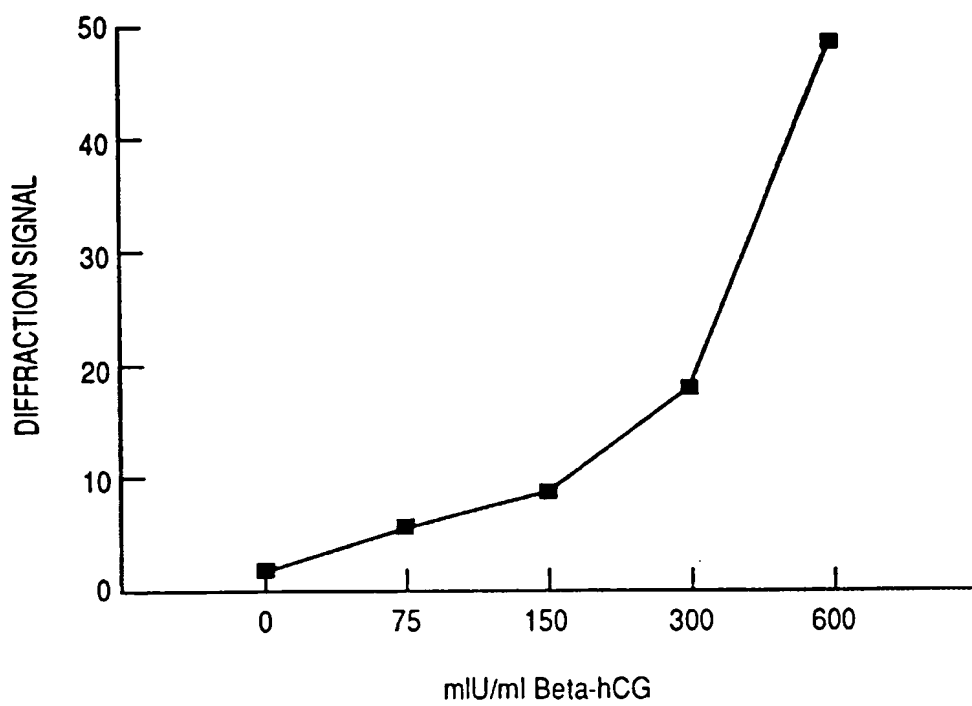


FIG. 5